## N(2-ACYLTHIO)PHENYLENAMIDES FROM BENZOTHIAZOLINES

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Enamides are interesting key intermediates in organic synthesis. Their preparation has been generally accomplished by acylation of imines  $^1$ , isomerization of allylamides  $^2$  and by elimination of methanol from  $\alpha$ -methoxylated amides  $^3$ .

We report now a new way which allows the synthesis of the enamides  $\underline{3}$  starting from easily available benzothiazolines  $\underline{1}$  and carboxylic anhydrides. The compounds  $\underline{3}$  are obtained in good yields according to the synthetic scheme below reported. The structures of the compounds  $\underline{3}$  were established by a combination of spectral and chemical evidences. The  ${}^1H$  nmr spectra revealed the presence of the vinylic proton in the region 4.95- $6.80~\delta$ . The hydrolysis of the compound  $\underline{3}$  yielded quantitatively the ketone  $\underline{4}$  and the amidethioester  $\underline{5}$ .

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